HEB SOLUTIONS SUNSCREEN ULTRA 30- avobenzone, homosalate, octisalate, octocrylene, oxybenzone lotion
H-E-B

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

HEB Solutions Sunscreen Ultra 30 Lotion

Active ingredients

Avobenzone 2%, Homosalate 7%, Octisalate 5%, Octocrylene 3%, Oxybenzone 4%

Purpose

Sunscreen

Uses

- Helps prevent sunburn
- If used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early aging caused by the sun

Warnings

For external use only

Do not use • on damaged or broken skin.

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if • rash occurs.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure.
- reapply:
- after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- children under 6 months of age: Ask a doctor
- **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:

- limit time in the sun especially from 10 a.m. – 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses

Other information

- protect the product in this container from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

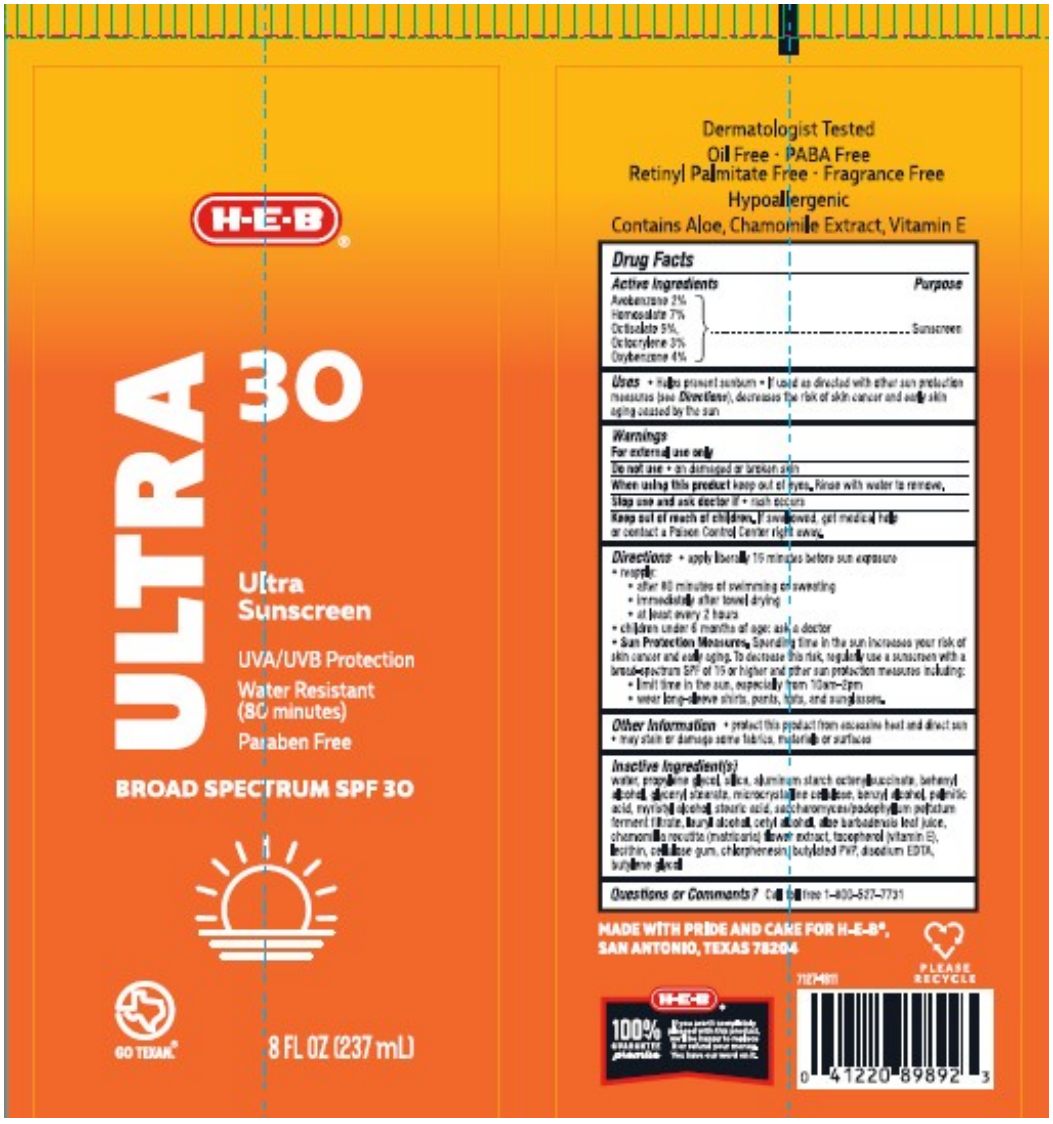
Inactive ingredients

water, propylene glycol, silica, aluminum starch octenylsuccinate, behenyl alcohol, glyceryl stearate, microcrystalline cellulose, benzyl alcohol, palmitic acid, myristyl alcohol, stearic acid, saccharomyces/podophyllum peltatum ferment filtrate, lauryl alcohol, cetyl alcohol, aloe barbadensis leaf juice, chamomilla recutita (matricaria) flower extract, tocopherol (vitamin E), lecithin, cellulose gum, chlorphenesin, butylated PVP, disodium EDTA, butylene glycol

HEB Solutions Sunscreen Ultra 30 Lotion

8 FL OZ (237)

NDC 37808-941-12



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Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-941
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	40 mg in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	20 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	30 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DOCOSANOL (UNII: 9G1OE216XY)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	
PALMITIC ACID (UNII: 2V16EO95H1)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
TOCOPHEROL (UNII: R0ZB2556P8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CHAMOMILE (UNII: FGL3685T2X)	
N-VINYLPYRROLIDINONE (UNII: 76H9G81541)	
YEAST (UNII: 3NY3SM6B8U)	
PODOPHYLLUM (UNII: 2S713A4VP3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-941-12	237 mL in 1 TUBE; Type 0: Not a Combination Product	02/06/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	02/06/2013	

Labeler - H-E-B (007924756)

Revised: 10/2022 H-E-B